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In the Claims

- 1. Cancel.
- 2. Cancel.
- 3. Cancel.
- 4. Cancel.
- 5. Cancel.
- Cancel. 6.
- 7.. Cancel.
- 8. Cancel.
- 9. Cancel.
- 10. Cancel.
- 11. Cancel.
- 12. Cancel.
- 13. Cancel.
- 14. Cancel.
- 15. Cancel.
- 16. Cancel.

17(Original) A compound of structural formula I:

I

or a pharmaceutically acceptable salt, enantiomer, diastereomer, pro drug or mixture thereof, wherein

X is (CH2)n, O or S;

Y represents
$$(C(R^b)_2)_n$$
, triple bond, R^b or $R^b \to R^b$

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Ar₂ independently represent (CH₂)_mC₆₋₁₀aryl, (CH₂)_mC₅₋₁₀heteroaryl, (CH₂)_mC₃₋₁₀ heterocycloalkyl, (CH₂)_mC₃₋₈ cycloalkyl said cycloalkyl, heterocycloalkyl, aryl or heteroaryl unsubstituted or substituted with 1-3 groups of Ra-

Ra represents C1-6 alkoxy, C1-6 alkyl, CF3, nitro, amino, cyano, C1-6 alkylamino, or halogen;

Rb independently represents H, halogen, C1-6 alkyl, C3-6 cylcoalkyl or

--- represents a double or single bond;

n represents 0-4; and

m represents 0-8.

18(Original). The compound according to claim 17 wherein X and Y are $(CH_2)_n$, --- represents a double bond; and Ar₂ is phenyl.

19(Original). The compound according to claim 18 wherein X is (CH₂)_n and n is 1 and Y is $(CH_2)_n$ and n is 3.

20 Cancel.

21(Currently Amended) A pharmaceutical composition for treating ocular hypertension or glaucoma comprising a therapeutically effective amount of a compound of formula I of claim 17 as defined in any one of claims 1 to 10, or a pharmaceutically acceptable salt, enantiomer, diasteromer, prodrug, or mixture thereof, in association with a pharmaceutically acceptable carrier.

22 (Currently Amended) A composition according to claim 21 in a form for topical administration as a solution or suspension and further comprising a second active ingredient as defined in claim 12 or 13.

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- 23 Cancel.
- 24 Cancel.
- 25 Cancel.
- 26 Cancel.
- 27 (New) A composition according to claim 22 wherein a second active ingredient belonging to the group consisting of: b-adrenergic blocking agent, parasympatho-memetic agent, sympathomimetic agent, carbonic anhydrase inhibitor, Maxi-K channel blocker, a prostaglandin, hypotensive lipid, neuroprotectant, and 5-HT2 receptor agonist is added.
- 28 (New) A composition according to claim 27 wherein the βadrenergic blocking agent is timolol, betaxolol, levobetaxolol, carteolol, or levobunolol; the parasympathomimetic agent is pilocarpine; the sympathomimetic agent is epinephrine, brimonidine, iopidine, clonidine, or para-aminoclonidine, the carbonic anhydrase inhibitor is dorzolamide, acetazolamide, metazolamide or brinzolamide; the prostaglandin is latanoprost, travaprost, unoprostone, rescula, or \$1033, the hypotensive lipid is lumigan, the neuroprotectant is eliprodil, R-eliprodil or memantine; and the 5-HT2 receptor agonist is 1-(2-aminopropyl)-3methyl-1H-imdazol-6-ol fumarate or 2-(3-chloro-6-methoxy-indazol-1yl)-1-methyl-ethylamine.